all skin blemishes, and clear up disturbed skin; and 502(e)(2)—the label of the article failed to bear the common or usual name of each active ingredient.

DISPOSITION: 5-15-61. Consent—claimed by Scandia Sales Corp., and released under bond for relabeling.

6692. Rel-Ong. (F.D.C. No. 45729. S. No. 72-573 R.)

QUANTITY: 780 1/4-oz btls. at Whittier, Calif.

SHIPPED: Between 12-6-60 and 3-1-61, from Trilby, Fla., by Meche Mfg. Co. LABEL IN PART: (Btl.) "REL-ONG BY MECHE 1956 THE FINGERNAIL BEAUTIFIER Meche Mfg. Co., Trilby, Fla. \* \* \* ½ Fl. Oz. \* \* \* Rel-Ong is not a polish, not a plastic nail you brush on. \* \* Active ingredients: Iodide Tinctures, Distillate of Hamamelidaceae, blended with essential oils and alcohol added as a preservative. Warning: \* \* \* Antidote: \* \* \*

Directions:."

Accompanying Labeling: Display cards reading in part "Do Your Nails Break or Split? Are They Thin? Brittle? \* \* \* Meche's Rel-Ong the Fingernail Beautifier"; folders entitled "Here They Are! Long Natural Glamor Nails – They Can Be Yours"; letters entitled "Rel-Ong What? – How? – Why?"; and Food and Drug Administration "Notice of Inspection" forms (FD Form 482) imprinted "Here's Positive Proof."

LIBELED: 4-24-61, S. Dist. Calif.

CHARGE: 502(a)—when shipped, the labeling contained false and misleading representations that the article would prevent and correct weak, thin, or brittle nails; aid nail growth; make nails grow longer, stronger, and thicker; prevent chipping, peeling, and breaking of nails; and that the article had the approval of the Food and Drug Administration.

Disposition: 6-5-61. Default—destruction.

6693. Torulose tablets and powder, and JerOtene capsules. (F.D.C. No. 46053. S. Nos. 20–366/8 R.)

QUANTITY: 2 drums containing 59,000 tablets, 50 250-tablet btls., 14 500-tablet btls., and 1 1,000-tablet btl. of *Torulose tablets*; 46 1-lb. cans of *Torulose powder*; and 1 box containing 10,000 capsules of *JerOtene*, at Kalamazoo, Mich., in possession of Torrance Co.

SHIPPED: Torulose tablets, 7-3-61, from Chicago, Ill.; Torulose powder, 6-21-61, from Rhinelander, Wis.; and JerOtene capsules, 6-22-61, from South Whitley, Ind.

Label in Part: (Drum) "Private Formula No. P-29.241 \* \* \* Prepared for The Torrance Company. \* \* \* Code #60824 F \* \* \* Each tablet contains: Torula Yeast 7½ gr."; (btl.) "Torulose The King of Food Yeasts with the pleasing nut like flavor grown on the 'Sweet of the Spruce' Dosage: \* \* \* Manufactured for Jerico Laboratories Kalamazoo, Michigan 8338"; (can) "Instant Yeast . . . With a Nut Like Flavor . . . Torulose \* \* \* King of Food Yeasts \* \* \* Manufactured for Jerico Laboratories, Kalamazoo, Michigan \* \* \* Each Tablespoonful (8 grams) Contains: \* \* \* Compared with a typical Brewer's Yeast \* \* \* 60824"; (btl.) "JerOtene \* \* \* Completely Organic 5000 U.S.P. Vitamin A Units of Carotene. 10 Intl. Units Vitamin E Dosage \* \* \* Jerico Laboratories Kalamazoo, Michigan Designated for special dietary food purposes \* \* \* Control No. 94173 Code No. 100."

Accompanying Labeling: Leaflets entitled "For Radiant Health Use Torulose" and "JerOtene Organic Carotene A Capsules"; also additional bottle labels for Torulose tablets and JerOtene capsules.

RESULTS OF INVESTIGATION: The Torulose tablets and powder in the bottles and cans were shipped in bulk and repacked and labeled by the dealer and, in the ordinary course of the dealer's business, the JerOtene capsules in the box were to be repacked into bottles labeled as described above.

LIBELED: 6-29-61, W. Dist. Mich.; amended libel filed 7-7-61.

CHARGE: Torulose tablets and powder, 502(a)—while held for sale, the labeling contained false and misleading representations that the articles were adequate and effective to promote radiant health and to promote the development of champions.

JerOtene capsules, 502(a)—while held for sale, the labeling contained false and misleading representations that the article was adequate and effective to promote growth, appetite and digestion; maintain a healthy appearance of hair and fingernails; maintain healthy balance of epithelial and nerve tissues; and to prevent infection of the respiratory system.

The libel alleged also that the *Torulose tablets and powder* and another article were misbranded under the provisions of the law applicable to foods, as reported in notices of judgment on foods.

DISPOSITION: 7-11-61. Consent—claimed by the dealer and released under bond for relabeling.

6694. Sacro-Disc-30. (F.D.C. No. 44926. S. No. 4-563 R.)

QUANTITY: 346 100-tablet btls. at Baltimore, Md., in possession of Armisyl Products Co.

Shipped: 5-27-60, from Philadelphia, Pa.

LABEL IN PART: (Btl.) "Sacro-Disc-30 Aneurine HC1 100 mg. U.S.P. Dose: 3 Tablets Daily \* \* \* 12471 Distributed by Armisyl Products, 2806 W. Strathmore Ave. Baltimore 9, Md."

Accompanying Labeling: Brochures entitled "Suffering from Backache? Leg Pain?"

RESULTS OF INVESTIGATION: The article was repacked by the Barre Drug Co., Inc., Baltimore, Md., after shipment as described above. The bottles and labels for use in repacking the article were supplied by the Armisyl Products Co. The above-mentioned brochures were prepared locally for the Armisyl Products Co. Analysis showed that the article contained thiamine hydrochloride.

Libeled: 10-3-60, Dist. Md.

CHARGE: 502(a)—while held for sale, the name "Sacro-Disc-30" and statements in the accompanying labeling of the article represented and suggested that the article was an adequate and effective treatment for backaches, leg pains, slipped disc, pains in general, arthritis, rheumatism, stiff joints, sleeplessness, nervous disorders, such as memory defects, depression, and severe agitation, and that the article was essential both to fertility and growth, and important to nerves, appetite, and digestion, which name and statements were false and misleading since the article was not an adequate and effective treatment for such conditions and purposes and was not capable of fulfilling the promises of benefit made for it; and 502(e)(2)—